REQUEST FOR QUOTATION FOR GOODS AND SERVICES

PM

ONDERSTEPOORT BIOLOGICAL PRODUCTS LTD
PRIVATE BAG X7, ONDERSTEPOORT 0110

From: Supply Chain Department Date: Oct 13 2025 Tel: 012 522 1500 Fax: N/A Email: purchasing@obpvaccines.co.za	To: Supplier: Tel: Fax: Email:

Kindly provide the quotation for the following: RFQ/OBP207/2025/26

Compulsory Document Requirements	Yes/No
Provide proof of previous supply for the product being requested or similar in the past 24 months (stamped Invoices or delivery note or reference letters). must be a minimum of 2	
SOUTH AFRICAN BIDDERS: Must be registered on CSD (active status) and provide a CSD report not older than 2 months (using the RFQ closing date). INTERNATIONAL BIDDERS: Wishing to bid must request an SBD 1 from the Procurement department (purchasing@obpvaccines.co.za) document to accompany with bid application.	
SBD4 Bidders Disclosure - All suppliers MUST Complete, sign & submit the SBD4 declaration with their bid application.	

Evaluation of Price and Preference

All Bids will be evaluated on a points system based on weighted average score for Price and Preference as per Preferential Procurement Framework Act of 2000 (Act 5 of 2000).

Preference Point allocation – 80/20	
Price / Preference	**Weighting percentage**
Preference:	**20%**
Price:	**80 %**
Total must equal:	**100%**

OBP Onderstepoort Biological Products will award preference points as follows: Specific Goal	Points	Evidence required	Yes/No

Historically disadvantaged by unfair discrimination on the basis of Race	10	A valid BBBEE Certificate showing at least 51% black ownership	
Historically disadvantaged by unfair discrimination on the basis of Gender (women)	8	A valid BBBEE Certificate showing at least 30% women ownership	
Historically disadvantaged by unfair discrimination on the basis of disability	2	A doctor's note confirming disability, confirmation of disability from the Department of Labour, BEE certificate or equivalent confirmation.	
Total points	**20**		

NB: Please note that if any of the above requirements is not submitted with the quote it will be an immediate disqualification.

TO APPOINT A SUPPLIERTO PROVIDE THE FOLLOWING ITEM/S OR SERVICE AS PER SCOPE BELOW.

Quantity	Product/Item Code	Specification
5000 EACH	Inserts for RVF CLONE 13	See attached specifications
10000 EACH	Cartons for PULPY KIDNEY Alum 100 ml	See attached specifications
5000 EACH	Cartons for RVF CLONE 13	See attached

Requirements from the supplier (To be used to select the contractor)

- The following will be required from a new bidder during evaluation stage:

- Sample will be requested

- Be audited by our Quality Assurance department before being approved/awarded

Government Procurement: all quotations of goods and services are subject to the General conditions of Contract July 2010

Requirements from SCM department:

- All bidders MUST register their company (in advance) on the NEW OBP's E-Procurement portal, the link can be found on the official OBP website under supply chain.
- Once bidders account registration is approved by the OBP Supply Chain, login credentials will be supplied, whereby bidders will be able to login and apply for opportunities.
- All open opportunities will reflect on the portal for bidders to part take in.
- All required company documents, proposed submissions or additional requirements MUST be uploaded wit you bid application.
- Any additional questions or Queries can be directed via email (purchasing@obpvaccines.co.za) or telephone (012 522 1500), note NO SUBMISSIONS WILL BE ACCEPTED via EMAIL.
- OBP reserves the right to cancel or re-advertise RFQ's (Request for quotes).

BIDDER'S DISCLOSURE

1. PURPOSE OF THE FORM

Any person (natural or juristic) may make an offer or offers in terms of this invitation to bid. In line with the principles of transparency, accountability, impartiality, and ethics as enshrined in the Constitution of the Republic of South Africa and further expressed in various pieces of legislation, it is required for the bidder to make this declaration in respect of the details required hereunder.

Where a person/s are listed in the Register for Tender Defaulters and / or the List of Restricted Suppliers, that person will automatically be disqualified from the bid process.

2. Bidder's declaration

2.1 Is the bidder, or any of its directors / trustees / shareholders / members / partners or any person having a controlling interest1 in the enterprise, employed by the state? **YES/NO**

2.1.1 If so, furnish particulars of the names, individual identity numbers, and, if applicable, state employee numbers of sole proprietor/ directors / trustees / shareholders / members/ partners or any person having a controlling interest in the enterprise, in table below.

Full Name	**dentity Number**	**Name of State institution**

2.2 Do you, or any person connected with the bidder, have a relationship with any person who is employed by the procuring institution? **YES/NO**

2.2.1 If so, furnish particulars:

……
……

2.3 Does the bidder or any of its directors / trustees / shareholders / members / partners or any person having a controlling interest in the enterprise have any interest in any other related enterprise whether or not they are bidding for this contract?

YES/NO

2.3.1 If so, furnish particulars:

……
……

3. DECLARATION

I, the undersigned, (name)……. in submitting the accompanying bid, do hereby make the following statements that I certify to

1 the power, by one person or a group of persons holding the majority of the equity of an enterprise, alternatively, the person/s having the deciding vote or power to influence or to direct the course and decisions of the enterprise.

be true and complete in every respect:

3.1 I have read, and I understand the contents of this disclosure.

3.2 I understand that the accompanying bid will be disqualified if this disclosure is found not to be true and complete in every respect.

3.3 The bidder has arrived at the accompanying bid independently from, and without consultation, communication, agreement, or arrangement with any competitor. However, communication between partners in a joint venture or consortium2 will not be construed as collusive bidding.

3.4 In addition, there have been no consultations, communications, agreements, or arrangements with any competitor regarding the quality, quantity, specifications, prices, including methods, factors or formulas used to calculate prices, market allocation, the intention or decision to submit or not to submit the bid, bidding with the intention not to win the bid and conditions or delivery particulars of the products or services to which this bid invitation relates.

3.5 The terms of the accompanying bid have not been, and will not be, disclosed by the bidder, directly or indirectly, to any competitor, prior to the date and time of the official bid opening or of the awarding of the contract.

3.6 There have been no consultations, communications, agreements, or arrangements made by the bidder with any official of the procuring institution in relation to this procurement process prior to and during the bidding process except to provide clarification on the bid submitted where so required by the institution; and the bidder was not involved in the drafting of the specifications or terms of reference for this bid.

3.7 I am aware that, in addition and without prejudice to any other remedy provided to combat any restrictive practices related to bids and contracts, bids that are suspicious will be reported to the Competition Commission for investigation and possible imposition of administrative penalties in terms of section 59 of the Competition Act No 89 of 1998 and or may be reported to the National Prosecuting Authority (NPA) for criminal investigation and or may be restricted from conducting business with the public sector for a period not exceeding ten (10) years in terms of the Prevention and Combating of Corrupt Activities Act No 12 of 2004 or any other applicable legislation.

I CERTIFY THAT THE INFORMATION FURNISHED IN PARAGRAPHS 1, 2 and 3 ABOVE IS CORRECT.
I ACCEPT THAT THE STATE MAY REJECT THE BID OR ACT AGAINST ME IN TERMS OF PARAGRAPH 6 OF PFMA SCM INSTRUCTION 03 OF 2021/22 ON PREVENTING AND COMBATING ABUSE IN THE SUPPLY CHAIN MANAGEMENT SYSTEM SHOULD THIS DECLARATION PROVE TO BE FALSE.

……..	…….
Signature	Date
…….	…….
Position	Name of bidder

Terms and Conditions:

- Submission should be no later than (**Oct 20 2025 15:00:00**)

2 Joint venture or Consortium means an association of persons for the purpose of combining their expertise, property, capital, efforts, skill and knowledge in an activity for the execution of a contract.

- Please indicate your offer validity and lead time: _____
- All prices must be VAT exclusive,(Vat vendor please indicate as such) if no indication, prices will be evaluated as exclusive.
- Quotation must be on a company letter head and ***strictly*** on a PDF format ***(Quotations sent on Word or Excel format will not be accepted***.)
- Supplier must register on or before any submission can be done , supplier number will be allocated to supplier.
- Submission and Quotations must be done online with all attachments required to be uploaded : any queries can be send to purchasing@obpvaccines.co.za
- **If no reply after 14 days of closing date your RFQ was unsuccessfully.**
- Please indicate if you are unable to quote and state the reason why
- Please note that fluctuations in the exchange rate (where applicable) will not be for the account of OBP.
- *Payment terms: 30 days after statement*
- *Bidders must be registered on CSD (Central Supplier Data Base National Treasury) and be tax complaint*
- **Government Procurement: all quotations of goods and services are subject to the General conditions of Contract July 2010**

I agree that the offer herein shall remain binding upon me and open for acceptance by OBP during the validity period indicated.

_____ _____

Signature Date

ENTEROTOXAEMIA ALUM VACCINE
100 ML DISPLAY CARTON
SUPPLIER SPECIFICATIONS
ITEM CODE : DC1231

Changes in future would be in **Bold** and *Italics.*

1. SPECIFICATION DETAILS

	REQUIREMENTS	SPECIFICATIONS
1	**Carton**	
1.1	Carton material	SBS (Cellulose) Solid bleach sulphate (Virgin board.) (Cellulose & ground wood pulp or wastepaper)
1.2	Carton material of construction	320 – 340 g/m²
1.3	Carton size	54 mm (L) x 54mm (W) (± 0,25 mm) x 92.5 mm (H) (± 0,25 mm), (Inner dimensions). (See ***drawing*** for full details of carton size)
1.4	Board colour	Both sides of the board must be white.
1.5	Direction of Grain	Direction of grain to be horizontal. (As per drawing)
1.6	All Carton flaps	Pre-cut is added on creasing line as per drawing. (See ***drawing***)
1.7	Carton stiffness	171TB 28 mNm longitudinal 67 TB 11 mNm cross
1.8	Varnish used	Note: Matt finish but no varnish / laminating on bottom flap for printing purposes. (See PRINTING AREA indicated on ***drawing***).
1.9	Adhesive used	26 AE or Equivalent.
1.10	Pre-breaking crease line	Pre-break carton at 140 to 180 degree and flatten again prior to making the longitudinal gluing seams.
1.11	Bar coding	Product bar code to be affixed on the top and bottom of the tucked flaps. The tucked flaps must be unpainted and unprinted. (See ***drawing***).
1.12	Bar code	12311
1.13	Top of the tucking flap	To be perforated for gluing and must be unprinted and unpainted. (See ***drawing***).
1.14	Bottom side flaps	To be perforated for gluing and must be unprinted and unpainted. (See ***drawing***).
2	**Printing**	
2.1	Pantone colours to comply to design specifications for each product	Pantone: 583 C, 5405 C, 7544 C, 7491 C.
2.2	Namibian Registration number	V01/24.4/157
2.3	Supplier identification	The supplier company identification must be indicated on the Display Carton.
2.4	Display carton reference, edition	The DC reference number and batch number must be

	and batch number on the bottom flap	printed on the bottom flap on the top left hand corner and the edition number must be printed on the bottom flap on the bottom right hand corner. Space must be provided for OBP to print the batch number and expiry date in the PRINTING AREA. (See ***drawing***).

3	**Die**	
3.1	Perforated ruling	The die must include special perforating ruling on the creases to assist machine packaging. (See ***drawing***).
3.2	Pre–Cut Crease line	The die must include pre-cuts on all the crease lines on all the flaps. (See ***drawing).***
4	**Special requirements**	
4.1	Gluing	No excess glue leakage must be present on the gluing tab of the carton.
4.2	Format of delivery and delivery	Die cut, creased, glued and delivered flat in bundles of 50, wrapped in paper. No rubber bands.
4.3	Flatened Box	Glue flap must be in center and the face up must be on the right side
4.4	Packed shipper requirements	The shipper must have a label with: • the Supplier name; • Supplier address; • Box / product name; • order number; • the quantity; and • The shipper must have a printed sample of the box attached to the outside as a quick reference.
5	**OBP acceptance criteria**	
5.1	Certificate of Conformance (C of C)	All consignments must have a Certificate of Conformance (C of C) containing the following: • Must be on company letter head; • State the batch number; • Name of carton printed and DC reference number; • Provide specifications and suppliers results of in house test of product; • Must indicate compliance with OBP's specification requirements and use same terminology; and • Results of supplier in-process tests.
5.2	Quality Assurance approval	• Before initial printing an uncut draw sheet with correct colours to be supplied to OBP for approval. • First print copies on final board must be supplied and approved by QA at OBP. • Mock-up box to be supplied and approved by OBP prior to printing.

ENTEROTOXAEMIA ALUM VACCINE
100 ML DISPLAY CARTON
SUPPLIER SPECIFICATIONS
ITEM CODE : DC1231

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100 ML DISPLAY CARTON
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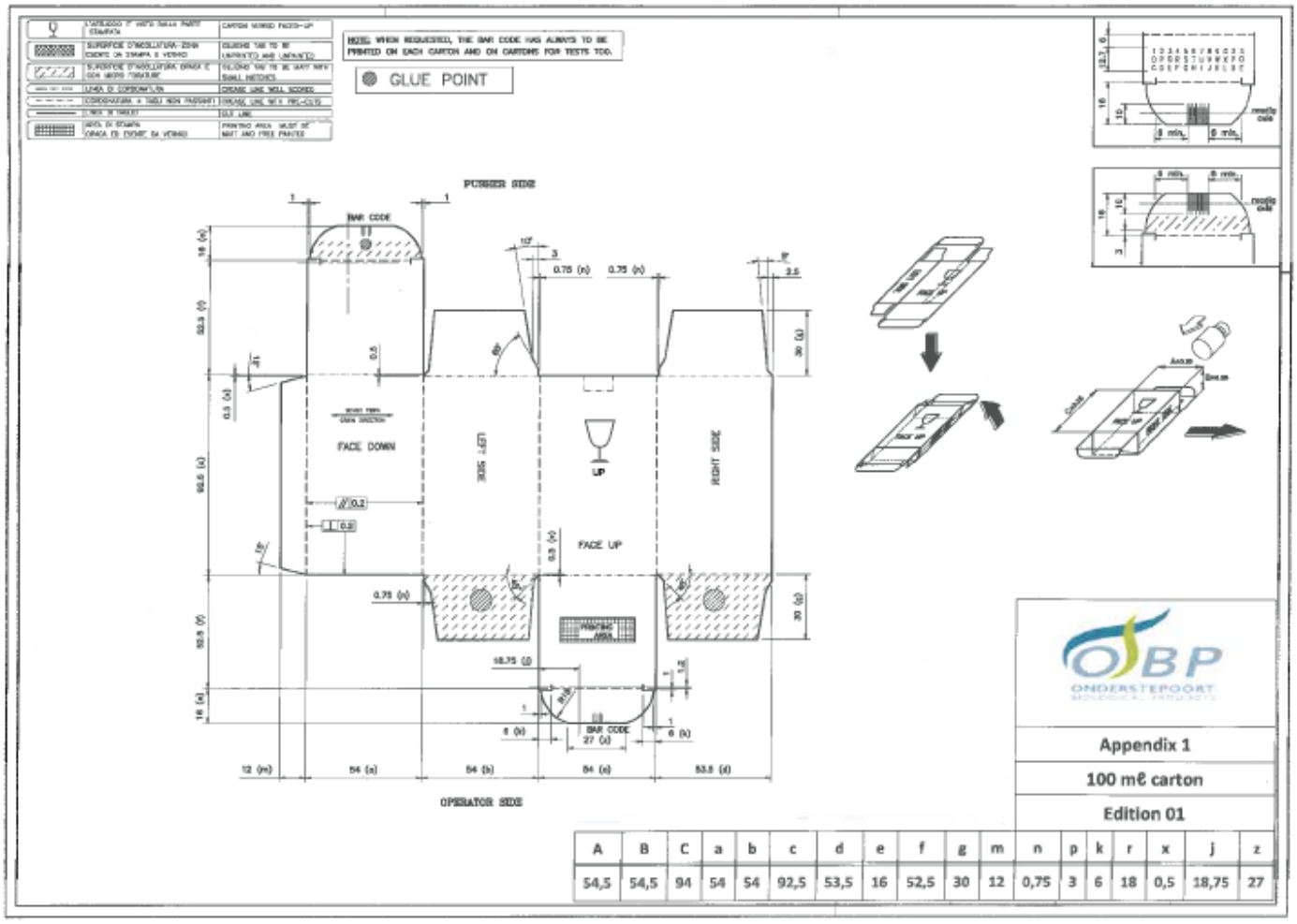
2. DRAWING



A	B	C	a	b	c	d	e	f	g	m	n	p	k	r	x	j	z
54,5	54,5	94	54	54	92,5	53,5	16	52,5	30	12	0,75	3	6	18	0,5	18,75	27

ENTEROTOXAEMIA ALUM VACCINE
100 ML DISPLAY CARTON
SUPPLIER SPECIFICATIONS
ITEM CODE : DC1231

3. ARTWORK

Author: Samuel Tshabangu Role: Packaging Manager Signature: Date:	Reviewed by: Raynard McDonald Role: QC Manager Signature: Date:	Approved by: Soveena Harriepersadh Role: QA Manager Signature: Date:

RIFT VALLEY FEVER CLONE 13 VACCINE
PACKAGE INSERT
SUPPLIER SPECIFICATIONS
ITEM CODE : PI 2213

OBP VP

Please note this is a new format. Changes in future would be in **Bold** and *Italics.*

1. Specification Details

REQUIREMENTS	SPECIFICATIONS
1. Material of construction	60 gsm Typek Bond.
2. Package insert dimensions	190 mm (L) x 130mm (W) (± 1,0 mm)
3. Package insert dimensions when folded	32 mm (L) X 130 mm (W) (± 1,0 mm)
4. Number of folds	3 fold position
5. Fiber direction	Fiber direction must be as per ***drawing***.
6. Thickness	55 micron (52 - 60 micron)
7. Paper colour	Surface is uncoated and white in colour
8. Colour	Process Black
9. Bar coding	Product bar code to be affixed on the top of the packaging insert as per ***drawing***.
10. Bar code	
11. Namibian Registration number	V10/24.4/958
12. Suppliers identification	The supplier company identification must be indicated on the bottom of the packaging insert.
13. Packaging Insert reference, and edition number	The PI reference number and edition number must be printed on the bottom of the packaging Insert on both sides.
14. Packing requirements	Each bundle of package inserts consists of ***300*** leaflets that is wrapped with paper finally packed into a shipper.
15. Inner box	***The inner boxes of leaflets must have labels with supplier name, insert package name, batch details and the quantity of the leaflets inside the inner box. The leaflets must be packed OR grouped into 50's inside the inner boxes.***
16. Packed shipper requirements	The shipper must have a label with • the Supplier name; • Supplier address; • Packaging insert / product name; • order number;

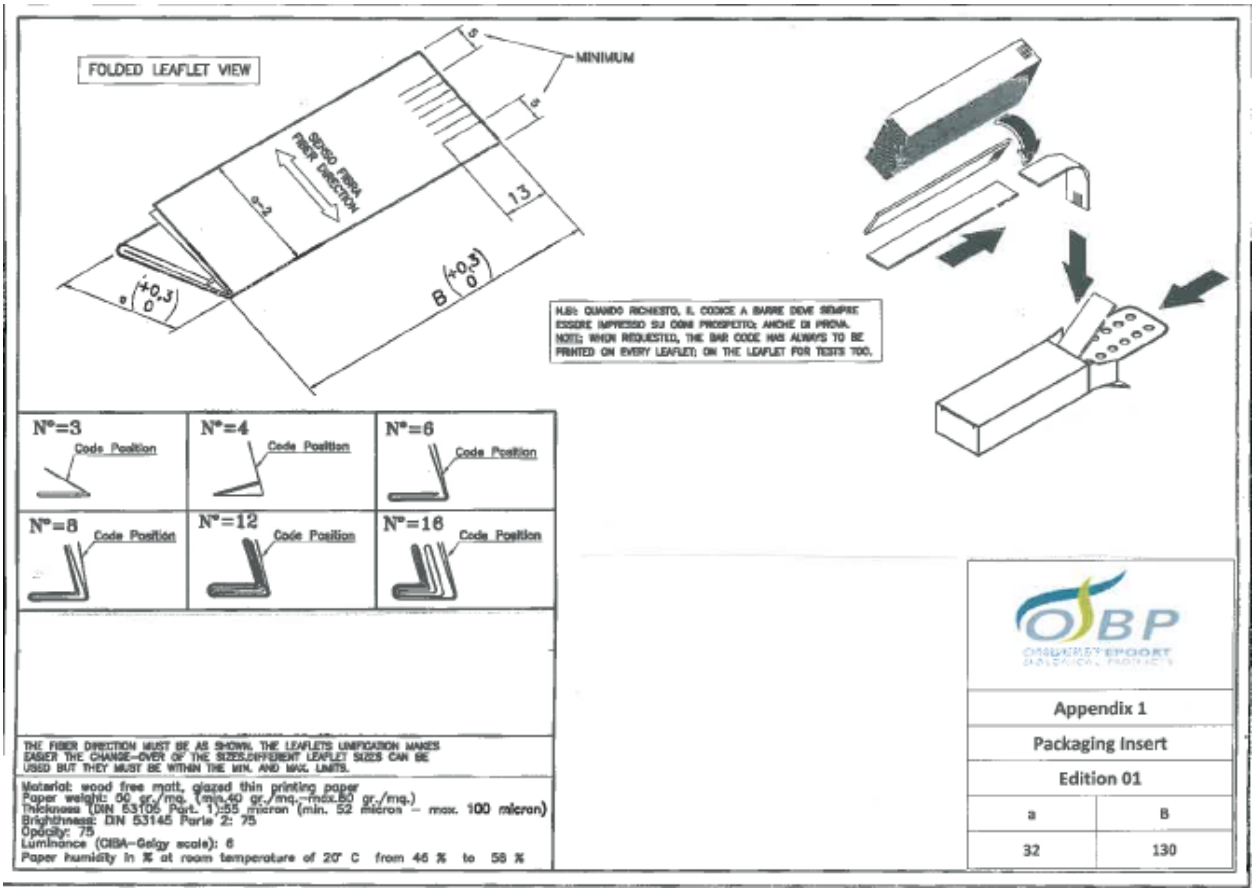
Author: Samuel Tshabangu Role: Packaging Manager Signature: Date:	Reviewed by: Raynard McDonald Role: QC Manager Signature: Date:	Approved by: Soveena Harriepersadh Role: QA Manager Signature: Date:

	• batch number • the quantity; and The shipper must have a printed sample of the Packaging insert attached to the outside of the box as a quick reference
17. Certificate of Conformance (C of C)	All consignments must have a Certificate of Conformance (C of C) containing the following: • Must be on company letter head; • State the batch number; • Name of packaging insert printed and Packaging insert reference number; • Provide specifications and suppliers results of in house test of product; • Must indicate compliance with OBP's specification requirements and use same terminology; and • Results of supplier in-process tests.
18. Qualiity Assurance Approval	• Before initial printing the supplier must provide OBP Q.A department a transparent template of the printing positive to be used for proof reading

RIFT VALLEY FEVER CLONE 13 VACCINE
PACKAGE INSERT
SUPPLIER SPECIFICATIONS
ITEM CODE : PI 2213

DRAWING

RIFT VALLEY FEVER CLONE 13 VACCINE
PACKAGE INSERT
SUPPLIER SPECIFICATIONS
ITEM CODE : PI 2213



Author: Samuel Tshabangu Role: Packaging Manager Signature: Date:	Reviewed by: Raynard McDonald Role: QC Manager Signature: Date:	Approved by: Soveena Harriepersadh Role: QA Manager Signature: Date:

1. ARTWORK

RIFT VALLEY FEVER CLONE 13 VACCINE
PACKAGE INSERT
SUPPLIER SPECIFICATIONS
ITEM CODE : PI 2213

ONDERSTEPOORT
BIOLOGICAL PRODUCTS

For animal use only

RVF CLONE 13
VACCINE FOR CATTLE, SHEEP AND GOATS

Reg. No. G 3876 (Act 36/1947) Namibia: V10 /24.4/958

Freeze-dried, live attenuated Rift Valley Fever virus (Clone 13 strain) for the immunisation of cattle, sheep and goats against Rift Valley Fever.

Store the vaccine at a temperature of 4 ˚C to 8 ˚C, in a refrigerator. Do not use the vaccine after the expiry date printed on the bottle.

RECOMMENDATIONS FOR USE
Rift Valley Fever is a mosquito-borne disease which occurs sporadically in South Africa especially during seasons of high rainfall. During outbreaks, the disease is prevalent in late summer to autumn (February - March). Animals should therefore be immunised in early summer to prevent infection. Young animals can be immunised with safety from 2 months of age, unless their mothers were vaccinated in which case they should be vaccinated at 6 months of age. **The vaccine has been shown to be safe for use in pregnant animals**. Vaccinate animals annually to ensure optimal immunity.

WARNINGS
Do not slaughter animals for human consumption within 7 days of vaccination. Vaccinate healthy animals only. Keep out of reach of children, uninformed persons and animals. Safe for use in pregnant animals. Although this product has been extensively tested under a large variety of conditions, failure thereof may ensue as a result of a wide range of reasons. If this is suspected, seek veterinary advice and notify the registration holder.

DIRECTIONS FOR USE
Use only as directed.
Sterilise syringes and needles by boiling in water for at least 15 minutes. Do not use disinfectants or methylated spirits for sterilising either needles or syringes. The active ingredient of the vaccine is in the form of powder or pellet in a small bottle. By means of a sterile syringe, transfer 5 ml diluent to the bottle containing freeze-dried vaccine. Mix thoroughly until the powder is dissolved. Transfer the suspension back to the remaining diluent and again mix well. The vaccine is now ready for use and it must be injected without delay. Avoid exposure to high temperatures and direct sunlight during inoculation. Shake the bottle before filling the syringe. Use a separate needle for each animal, particularly during outbreaks of the disease. After re-suspension of the vaccine, animals should be vaccinated within 2 hours.

DOSAGE:
Cattle, Sheep and Goats: **1 ml subcutaneously**

EFFECTS OF THE VACCINE
A slight febrile reaction may occur on the second to fourth day following inoculation but subsides rapidly. Most animals will be immune three weeks after vaccinations, although this cannot be guaranteed. Annual vaccination is recommended.

PACKING
Available in bottles of 100 doses.

Registration holder:
Onderstepoort Biological Products (Ltd), Co. Reg. No. 2000/022686/06
Private Bag X07, Onderstepoort, 0110. Tel: +27 (0) 12 522 1500, Fax: +27 (0) 12 522 1591

P22130

Lomar Printers Made in South Africa Edition 5

RIFT VALLEY FEVER CLONE 13 VACCINE
PACKAGE INSERT
SUPPLIER SPECIFICATIONS
ITEM CODE : PI 2213

Slegs vir dieregebruik

RVF CLONE 13
ENTSTOF VIR BEESTE, SKAPE EN BOKKE

Reg Nr. G 3876 (Wet 36/1947) Namibie: V10 /24.4/958

Lewende, verswakte Slenkdalkoorsvirus (Kloon 13 stam) in gevriesdroogde vorm vir die immunisering van beeste, skape en bokke teen Slenkdalkoors.

Bewaar die entstof by 'n temperatuur van 4 °C tot 8 °C, in 'n yskas. Moet nie die entstof na die vervaldatum wat op die bottel gedruk is gebruik nie.

AANBEVELINGS VIR GEBRUIK
Slenkdalkoors is 'n muskietoorgedraagde virussiekte wat sporadies in Suid-Afrika voorkom, veral tydens hoë reënval seisoene. Tydens uitbrake kom die siekte voor in laat somer tot herfs (Februarie-Maart). Diere moet dus vroeg in die somer geënt word om die siekte te voorkom. Jong diere kan met veiligheid vanaf 2 maande ouderdom geënt word, mits hul moeders nie geënt was nie in welke geval inenting op 6 maande ouderdom moet plaasvind. **Die entstof is veilig vir gebruik in dragtige diere**. Jaarlikse inenting word aanbeveel.

WAARSKUWINGS
Moet nie diere binne 7 dae na inenting vir menslike verbruik slag nie. Slegs gesonde diere moet ingeënt word. Hou buite die bereik van kinders, oningeligte persone en diere. Veilig vir gebruik in dragtige diere. Alhoewel hierdie produk breedvoerig onder 'n wye verskeidenheid van toestande getoets is, mag dit faal as gevolg van verskeie redes. Indien dit vermoed word, raadpleeg 'n veearts en verwittig die registrasiehouer.

GEBRUIKSAANWYSINGS
Gebruik slegs soos voorgeskryf.
Steriliseer spuite en naalde deur dit vir ten minste 15 minute in water te kook. Moet nie ontsmettingsmiddels of brandspiritus gebruik om spuite en naalde te steriliseer nie. Die aktiewe bestanddeel van die entstof is in die vorm van 'n poeier of klont in 'n klein botteltjie. Dra 5 ml verdunningsvloeistof met 'n steriele spuit oor na die botteltjie met gedroogde entstof en meng goed totdat die poeier opgelos is. Dra die opgeloste entstof terug na die oorgeblewe verdunningsvloeistof en meng weer goed. Die entstof is nou gereed vir gebruik en moet dadelik ingespuit word. Vermy blootstelling aan hoë temperature en direkte sonlig gedurende inenting. Skud die bottel goed voor gebruik. Gebruik 'n afsonderlike naald vir elke dier, veral tydens Slenkdalkoors uitbrake. Nadat die entstof opgelos is moet diere binne 2 ure ingespuit word.

DOSIS:
Beeste, Skape en Bokke: **1 ml onderhuids**

UITWERKING VAN DIE ENTSTOF
'n Ligte koorsreaksie wat gou verdwyn mag vanaf die tweede tot vierde dag na inenting voorkom en na drie weke behoort die diere ten volle beskerm te wees, alhoewel dit nie gewaarborg kan word nie. Jaarlikse enting word aanbeveel.

VERPAKKING
Beskikbaar in bottels van 100 dosisse.

Registrasiehouer:
Onderstepoort Biologiese Produkte (Bpk), Mpy. Reg. Nr. 2000/022686/06
Privaatsak X07, Onderstepoort, 0110. Tel: +27 (0) 12 522 1500, Faks: +27 (0) 12 522 1591

Vervaardig in Suid-Afrika Weergawe 5

RIFT VALLEY FEVER CLONE 13 VACCINE
100 ML COMBO DISPLAY CARTON
SUPPLIER SPECIFICATIONS
ITEM CODE : DC2213

OBP VP

Changes in future would be in **Bold** and *Italics.*

1. SPECIFICATION DETAILS

	REQUIREMENTS	SPECIFICATIONS
1	**Carton**	
1.1	Carton material	SBS (Cellulose) Solid bleach sulphate (Virgin board.) (Cellulose & ground wood pulp or wastepaper)
1.2	Carton material of construction	320 – 340 g/m²
1.3	Carton size	82 mm (L) x 54mm (W) (± 0,25 mm) x 92.5 mm (H) (± 0,25 mm) (Inner dimensions). (See drawing for full details of carton size)
1.4	Board colour	Both sides of the board must be white.
1.5	Direction of Grain	Direction of grain to be horizontal. (As per drawing)
1.6	All Carton flaps	Pre-cut is added on creasing line as per drawing. (See ***drawing***)
1.7	Carton stiffness	171TB 28 mNm longitudinal 67 TB 11 mNm cross
1.8	Varnish used	Machine varnish Note: Matt finish but no varnish / laminating on bottom flap for printing purposes. (See PRINTING AREA indicated on ***drawing***).
1.9	Adhesive used	26 AE or Equivalent.
1.10	Pre-breaking crease line	Pre-break carton at 140 to 180 degree and flatten again prior to making the longitudinal gluing seams.
1.11	Bar coding	Product bar code to be affixed on the top and bottom of the tucked flaps. The tucked flaps must be unpainted and unprinted. (See ***drawing***).
1.12	Bar code	22131
1.13	Top of the tucking flap	To be perforated for gluing and must be unprinted and unpainted. (See ***drawing***).
1.14	Bottom side flaps	To be perforated for gluing and must be unprinted and unpainted. (See ***drawing***).
2	**Printing**	
2.1	Pantone colours to comply to design specifications for each product	Pantone 583 C, 5405 C; 7544 C, 468 C.
2.2	Namibian Registration number	V10/24.4/958
2.3	Supplier identification	The supplier company identification must be indicated on

		the Display Carton.
2.4	Display carton reference, edition and batch number on the bottom flap	The DC reference number and batch number must be printed on the bottom flap on the top left hand corner and the edition number must be printed on the bottom flap on the bottom right hand corner. Space must be provided for OBP to print the batch number and expiry date in the PRINTING AREA. (See ***drawing***).

3	**Die**	
3.1	Perforated ruling	The die must include special perforating ruling on the creases to assist machine packaging. (See ***drawing***).
3.2	Pre–Cut Crease line	The die must include pre-cuts on all the crease lines on all the flaps. (See ***drawing).***
4	**Special requirements**	
4.1	Gluing	No excess glue leakage must be present on the gluing tab of the carton.
4.2	Format of delivery and delivery	Die cut, creased, glued and delivered flat in bundles of 50, wrapped in paper. No rubber bands.
4.3	Flatened Box	Glue flap must be in center and the face up must be on the right side
4.4	Packed shipper requirements	The shipper must have a label with: • the Supplier name; • Supplier address; • Box / product name; • order number; • the quantity; and • The shipper must have a printed sample of the box attached to the outside as a quick reference.
5	**OBP acceptance criteria**	
5.1	Certificate of Conformance (C of C)	All consignments must have a Certificate of Conformance (C of C) containing the following: • Must be on company letter head; • State the batch number; • Name of carton printed and DC reference number; • Provide specifications and suppliers results of in house test of product; • Must indicate compliance with OBP's specification requirements and use same terminology; and • Results of supplier in-process tests.
5.2	Quality Assurance approval	• Before initial printing an uncut draw sheet with correct colours to be supplied to OBP for approval.

RIFT VALLEY FEVER CLONE 13 VACCINE
100 ML COMBO DISPLAY CARTON
SUPPLIER SPECIFICATIONS
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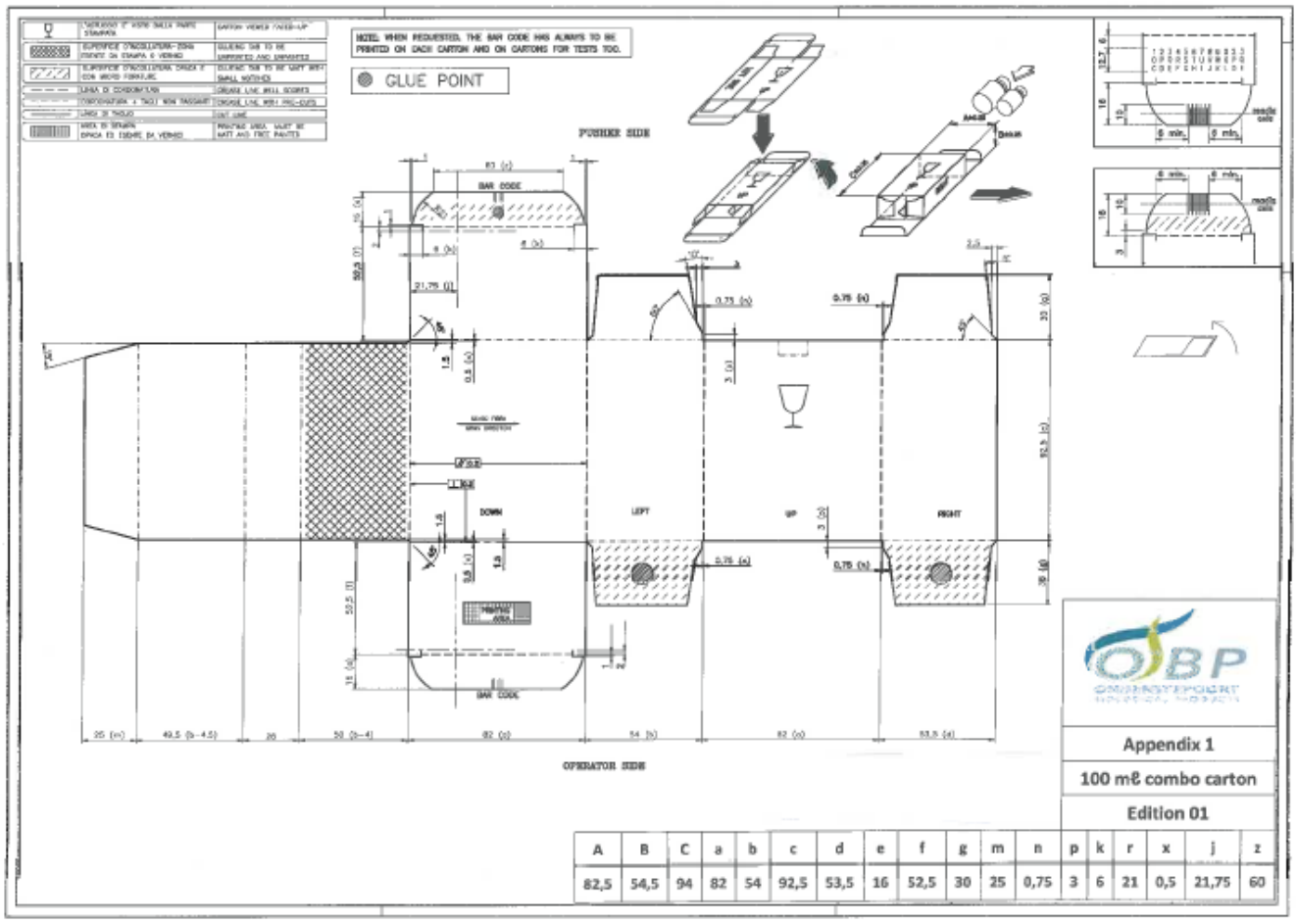
		• First print copies on final board must be supplied and approved by QA at OBP. • Mock-up box to be supplied and approved by OBP prior to printing.

RIFT VALLEY FEVER CLONE 13 VACCINE
100 ML COMBO DISPLAY CARTON
SUPPLIER SPECIFICATIONS
ITEM CODE : DC2213

2. DRAWING



A	B	C	a	b	c	d	e	f	g	m	n	p	k	r	x	j	z
82,5	54,5	94	82	54	92,5	53,5	16	52,5	30	25	0,75	3	6	21	0,5	21,75	60

RIFT VALLEY FEVER CLONE 13 VACCINE
100 ML COMBO DISPLAY CARTON
SUPPLIER SPECIFICATIONS
ITEM CODE : DC2213

3. ARTWORK

Author: Samuel Tshabangu Role: Packaging Manager Signature: Date:	Reviewed by: Raynard McDonald Role: QC Manager Signature: Date:	Approved by: Soveena Harriepersadh Role: QA Manager Signature: Date: